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Amendments to the claims begin on page 2.

Amendments to the specification begin on page 5.

Remarks begin on page 11.

Amendments to the claims:

1. (Withdrawn) A method for increasing the specific activity of a glycosyl hydrolase on a substrate, comprising replacing a hydrophobic surface binding amino acid of the hydrolase with a positively charged amino acid, to provide a mutant glycosyl hydrolase.

2. (Withdrawn) The method of claim 1, wherein the hydrophobic surface binding amino acid includes tryptophan or tyrosine and the positively charged amino acid is arginine.

3. (Currently Amended) A method for increasing the specific activity of a mutated glycosyl hydrolase on a substrate relative to an unmutated form of the glycosyl hydrolase, comprising replacing an active site associated glycosyl-stabilizing amino acid of the hydrolase with an amino acid, the replacing amino acid binding not strongly retarding cellobiose less tightly than the glycosyl-stabilizing amino acid from leaving the active site to provide a mutant glycosyl hydrolase.

4. (Currently Amended) The method of claim 3, wherein the glycosyl-stabilizing amino acid comprises tyrosine-3 and the replacing amino acid comprises glycine.

5. (Currently Amended) The methods of claims 1 and 3, wherein the replacing step comprises replacing by site-directed-mutagenesis.

6. (Currently Amended) The methods of claims 1 and 3, wherein the mutant glycosyl hydrolase comprises a mutant EI endoglucanase.

7. (Currently Amended) The methods of claims 1 and 3, wherein the mutant glycosyl hydrolase comprises SEQ ID NO: 2-Y245G, SEQ ID NO: 3 Y42R, SEQ ID NO: 4 W82R, or a mixture thereof.

8. (Currently Amended) The methods of claims 1 and 3, wherein the substrate comprises pretreated biomass.

9. (Withdrawn) A mutant glycosyl hydrolase having enhanced catalytic activity, said mutant glycosyl hydrolase comprising an amino having a positively charged amino acid at a position occupied by a hydrophobic surface binding amino acid in a wild-type

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glycosyl hydrolase amino acid sequence, wherein said mutant glycosyl hydrolase has an enhanced catalytic activity of 10% to 50% compared to catalytic activity of the wild-type glycosyl hydrolase.

10. (Withdrawn) The mutant glycosyl hydrolase of claim 9 further defined as a cellulase.

11. (Withdrawn) The mutant glycosyl hydrolase of claim 9 further defined as Y245G.

12. (Withdrawn) The mutant ~~glyeessy~~ glycosyl hydrolase of claim 9 further defined as Y42R.

13. (Withdrawn) The mutant glycosyl hydrolase of claim 9 further defined as a mannanase.

14. (Withdrawn) The mutant glycosyl hydrolase of claim 9 further defined as comprising W82R.

15 (Withdrawn) A method for converting a biomass into ethanol comprising: a. mixing a composition comprising biomass with a mutant glycosyl hydrolase having enhanced catalytic activity over a wild-type glycosyl hydrolase to provide a soluble fermentable sugar preparation; and b. fermenting said soluble fermentable sugar preparation to provide a composition comprising ethanol.

16. (Withdrawn) The method of claim 15 wherein the biomass is a cellulosic biomass.

17. (Withdrawn) The method of claim 15 wherein the mutant glycosyl hydrolase is Y245G.

18. (Withdrawn) The method of claim 15 wherein the mutant glycosyl hydrolase is Y82R.

19. (Withdrawn) The method of claim 15 wherein the mutant glycosyl hydrolase is W42R.

20. (Withdrawn) The method of claim 15 wherein the mutant glycosyl hydrolase comprises Y245G, Y82R, or W42R.

21. (Withdrawn) The method of claim 15 wherein the biomass is further admixed with a glycohydrolase.

22-25. (Cancelled)

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26. (Withdrawn) A method for increasing the specific activity of a hydrolytic depolymerizing enzyme, comprising replacing an extended-active site residue that binds strongly to the leaving group with another that binds much less strongly to the leaving group.

27. (Withdrawn) A method for increasing the specific activity of a glycosyl hydrolase on a pretreated biomass substrate, comprising:

replacing, via site directed mutagenesis, a hydrophobic surface binding amino acid of the hydrolase with a positively charged amino acid arginine, the hydrophobic surface binding amino acid of the hydrolase selected from the group consisting of tryptophan and tyrosine, the hydrolase selected from the group consisting of an EI endoglucanase, Y245G, Y42R, W82R, and a mixture thereof; and

replacing, via site directed mutagenesis, an active site associated glycosyl-stabilizing amino acid of the hydrolase including tyrosine with an amino acid including glycine, the glycine not strongly retarding cellobiose from leaving the active site,

to produce a mutant glycosyl hydrolase having enhanced catalytic activity, the mutant glycosyl hydrolase comprising an amino having a positively charged amino acid at a position occupied by a hydrophobic surface binding amino acid in a wild-type glycosyl hydrolase amino acid sequence, wherein said mutant glycosyl hydrolase has an enhanced catalytic activity of 10% to 50% compared to catalytic activity of the wild-type glycosyl hydrolase.

28. (Withdrawn) The method of claim 27, further comprising steps of mixing a composition comprising biomass with the mutant glycosyl hydrolase having enhanced catalytic activity over a wild-type glycosyl hydrolase to provide a soluble fermentable sugar preparation; and

fermenting the soluble fermentable sugar preparation to provide a composition comprising ethanol.

29. (New) A method for increasing the specific activity of an EI endoglucanase or a structural analog thereof on a biomass, comprising replacing, by site-directed-mutagenesis, an active site associated glycosyl-stabilizing amino acid of the endoglucanase with an amino acid, the replacing amino acid binding cellobiose less tightly than the glycosyl-stabilizing amino acid to provide a mutant endoglucanase.

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30. (New) The method of claim 29, wherein the glycosyl-stabilizing amino acid comprises tyrosine and the replacing amino acid comprises glycine.

31. (New) The method of claim 29, wherein the mutant endoglucanase comprises SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, or a mixture thereof.

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On pages 28-30 of the Specification, please replace the paragraphs including all of Example 7 in its entirety beginning at "Example 7" and ending prior to "BIBLIOGRAPHY" with the following paragraphs. These paragraphs include paragraphs 72-80 in the published application. There are no deletions in the text below, and additions are indicated by double underlining because portions of the inserted text contain native underlining.

EXAMPLE 7**Sequence Information**

The following table provides sequence data referenced throughout the present specification.

Nucleic acid sequence for EI endoglucanase

5'-

GCGGGCGGCGGCTATTGGCACACGAGCGGCCGGGAGATCCTGGACGCGAAC
AACGTGCCGGTACGGATCGCCGGCATCAACTGGTTTGGGTTTCGAAACCTGCA
ATTACGTCGTGCACGGTCTCTGGTCACGCGACTACCGCAGCATGCTCGACCA
GATAAAGTCGCTCGGCTACAACA- CAATCCGGCTGCCGTACTCTGACGAC
ATTCTCAAGCCGGGCACCATGCCGAACAGCATCAATTTTACCAGATGAATC
AGGACCTGCAGGGTCTGACGTCCTTGAGGTCATGGACAAAATCGTCGCGTA
CGCCGGTCAGATCGGCCTGCGCATCATTCTTGACCGCCACCGACCGGATTGC
AGCGGGCAGTCGGCGCTGTGGTACACGAGCAGCGTCTCGGAG- GCTACGT
GGATTTCCGACCTGCAAGCGCTGGCGCAGCGCTACAAGGGAAACCCGACG-
GTCGTCGGCTTTGACTTGACACAACGAGCCGCATGACCCGGCCTGCTGGGGCT
GCGGCGATCCGAGCATCGACTGGCGATTGGCCGCGAGCGGGCCGGAAACG
CCGTGCTCTCG- GTGAATCCGAACCTGCTCATTTCGTCGAAGGTGTGCAGA
GCTACAACGGAGACTCCTACTGGTGGGGCGGCAACCTGCAAGGAGCCGGCC
AGTACCCGGTCGTGCTGAACGTGCCGAACCGCCTGGTGTACTCGGCGCACGA
CTACGCGACGAGCGTCTACCCGCGAGACGTGGTTCAGCGATCCGACCTTCCCC
AACAACATGCCCCGGCATCTGGAACAAGAACTGGGGATACCTCTTCAATC

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AGAACATTGCACCGGTATGGCTGGGCGAATTCGGTACGACACTGCAATCCAC
GACCGACCAGACGTTGGCTGAAGACGCTCGTCCAGTACCTACGGCCGACCGC
GCAATACGGTGCGGACAGCTTCCAGTGGACCTTCTGGTCCTGGAACCCCGAT
TCCGGCGACACAGGAGGAATTCTCAAGGATGACTGGCAGACGGTCGACACA
GTAAAAGACGGCTATCTCGCGCCGATCAAGTCGTCGATTTTCGATCCTGTCTA
ATGAATCGCCTAGCAGTCAACCGTCCCCGTCGGTGTCGCCGTCTCCGTCGCCG
AGCCCGTCGGCGAGTCGGACGCCGACGCCTACTCCGACGCCGACAGCCAGCC
CGACGCCAACGCTGACCCCTACTGCTACGCCACGCCACGGCAAGCCCGAC
GCCGTCACCGACGGCAGCCTCCGGAGCCCGCTGCACCGCGAGTTACCAGGTC
AACAGCGATTGGGGCAATGGCTTCACGGTAACGGTGGCCGTGACAAATTCCG
-3' [SEQ ID NO. 1]

Amino acid sequence for EI endoglucanase

5-

AGGGYWHTSGREILDANNVPVRLAGINWFGFETCNVYVHGLWSRDYRSMLDQI
KSLGYNTIRLPYSDDILKPGTMPNSINFYQMNQDLQGLTSLQVMDKIVAYAGQIG
LRILDRHRPDCSGQSALWYTSSVSEATWISDLQALAQRYKGNPTVVGFDLHNEP
HDPACWGCSDPSIDWRLAAERAGNAVL SVNPNLLIFVEGVQSYNGDSY
WWGGNLQGAGQYPVVLNVPNRLVYSAHDYATSVYPQTWFSPTFPNNMPGIW
NKNWGYLFNQNIAPVWLGEFGTTLQSTTDQTLVQYLRPTAQYGADSFQW
TFWSWNPDSGDTGGILKDDWQTVDTVKDGYLE PIKSSIFDPVG-3' [SEQ ID NO.

2-

DNA sequence for Y245G Mutant with mutation site underlined.

5-

GCGGGCGGCGGCTATTGGCACACGAGCGGCCGGGAGATCCTGGACGCGAAC
AACGTGCCGGTACGGATCGCCGGCATCAACTGGTTTGGGTTCGAAACCTGCA
ATTACGTCGTGCACGGTCTCTGGTCACGCGACTACCGCAGCATGCTCGACCA
GATAAAGTCGCTCGGCTACAACA- CAATCCGGCTGCCGTACTCTGACGAC
ATTCTCAAGCCGGGCACCATGCCGAACAGCATCAATTTTACCAGATGAATC
AGGACCTGCAGGGTCTGACGTCCTTGCAGGTCATGGACAAAATCGTCGCGTA

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CGCCGGTCAGATCGGCCTGCGCATCATTCTTGACCGCCACCGACCGGATTGC
AGCGGGCAGTCGGCGCTGTGGTACACGAGCAGCGTCTCGGAGGCTACGT
GGATTTCCGACCTGCAAGCGCTGGCGCAGCGCTACAAGGGAAACCCGACG-
GTCGTCGGCTTTGACTTGCACAACGAGCCGCATGACCCGGCCTGCTGGGGCT
GCGGCGATCCGAGCATCGACTGGCGATTGGCCGCCGAGCGGGCCGGAAACG
CCGTGCTCTCG- GTGAATCCGAACCTGCTCATTTTCGTCGAAGGTGTGCAGA
GCTACAACGGGAGACTCCTACTGGTGGGGCGGCAACCTGCAAGGAGCCGGCC
AGTACCCGGTCGTGCTGAACGTGCCGAACCGCCTGGTGTACTCGGCGCACGA
CTACGCGACGAGCGTCGGCCCGCAGACGTGGTTTCAGCGATCCGACCTTCCCC
AACAACATGCCCGGCATCTGGAACAAGAACTGGGGATACCTCTTCAATCAGA
ACATTGCACCGGTATGGCTGGGCGAATTCGGTACGACACTGCAATCCACGAC
CGACCAGACGTGGCTGAAGACGCTCGTCCAGTACCTACGGCCGACCGCGCAA
TACGGTGCGGACAGCTTCCAGTGGACCTTCTGGTCCTGGAACCCCGATTCCG
GCGACACAGGAGGAATTCTCAAGGATGACTGGCAGACGGTCGACACAGTAA
AAGACGGCTATCTCGCGCCGATCAAGTCGTCGATTTTCGATCCTGTCTAATGA
ATCGCCTAGCAGTCAACCGTCCCCGTCGGTGTGCGCCGTCTCCGTGCGCCGAGCC
CGTCGGCGAGTCGGACGCCGACGCCTACTCCGACGCCGACAGCCAGCCCGAC
GCCAACGCTGACCCCTACTGCTACGCCACGCCACGGCAA
GCCCGACGCCGTCACCGACGGCAGCCTC-
CGGAGCCCGCTGCACCGCGAGTTACCAGGTCAACAGCGATTGGGGCAAT-3'

[SEQ ID NO. 2]

Translated amino acid sequence for Y245G mutation, with modification
underlined.

5'-

AGGGYWHTSGREILDANNVPVRIAGINWFGFETCNVYVHGLWSRDYRSM LDQI
KSLGYNTIRLPYSDDILKPGTMPNSINFYQMNQDLQGLTSLQVMDKIVAYAGQIG
LRILDRHRPDCSGQSALWYTSSVSEATWISDLQALAQRYKGNPTVVGF DLHNEP
HDPACWGCGDPSIDWRLAAERAGNAVL SVNPNLLIFVEGVQSYNGDSYWWGG
NLQGAGQYPVVLNVPNRLVYSAHDYATSVGPQTWFS DPTFPNNMPGIWNKNW

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GYLFNQNIAPVWLGEFGTTLQSTTDQTLVQYLRPTAQYGADSFQWTFWS
WNPDSGDTGGILKDDWQTVDTVKGDLA PIKSSIFDPV-3' [SEQ ID NO. 2]

DNA sequence for W42R Mutant with mutation site underlined

5'-

GCGGGCGGCGGCTATTGGCACACGAGCGGCCGGGAGATCCTGGACGCGAAC
AACGTGCCGGTACGGATCGCCGGCATCAACTGGTTTGGGTTCGAAACCTGCA
ATTACGTCGTGCACGGTCTCCGGTTCACGCGACTACCGCAGCATGCTCGACCA
GATAAAGTCGCTCGGCTACAACACAATCCGGCTGCCGTACTCTGACGAC
ATTCTCAAGCCGGGCACCATGCCGAACAGCATCAATTTTACCAGATGAATC
AGGACCTGCAGGGTCTGACGTCCTTGCAGGTCATGGACAAAATCGTCGCGTA
CGCCGGTTCAGATCGGCCTGCGCATCATTCTTGACCGCCACCGACCGGATTGC
AGCGGGCAGTCGGCGCTGTGGTACACGAGCAGCGTCTCGGAGGCTACGTGGA
TTTCCGACCTGCAAGCGCTGGCGCAGCGCTACAAGGGAAACCCGACGGTCGT
CGGCTTTGACTTGACAACGAGCCGCATGACCCGGCCTGCTGGGGCTGCGGC
GATCCGAGCATCGACTGGCGATTGGCCGGCGAGCGGGCCGGAAACGCCGTGC
TCTCGGTGAATCCGAACCTGCTCATTTCGTCGAAGGTGTGCAGAGCTACAAC
GGAGACTCCTACTGGTGGGGCGGCAACCTGCAAGGAGCCGGCCAGTACCCG
GTCGTGCTGAACGTGCCGAACCGCCTGGTGTACTCGGCGCACGACTACCGA
CGAGCGTCTACCCGCAGACGTGGTTCAGCGATCCGACCTTCCCCAACAACAT
GCCCCGCATCTGGAACAAGAACTGGGGATACCTCTTCAATCAGAACATTGCA
CCGGTATGGCTGGGCGAATTCGGTACGACACTGCAATCCACGACCGACCAGA
CGTGGCTGAAGACGCTCGTCCAGTACCTACGGCCGACCGCGCAATACGGTGC
GGACAGCTTCCAGTGGACCTTCTGGTCCTGGAACCCCGATTCCGGCGACACA
GGAGGAATTCTCAAGGATGACTGGCAGACGGTCGACACAGTAAAAGACGGC
TATCTCGCGCCGATCAAGTCGTCGATTTTCGATCCTGTCTAATGAATCGCCTA
GCAGTCAACCGTCCCCGTCGGTGTGCGCGTCTCCGTCGCCGAGCCCGTCGGC
GAGTCGGACGCCGACGCCTACTCCGACGCCGACAGCCAGCCCGACGCCAACG
CTGACCCCTACTGCTACGCCCACGCCCACGGCAAGCCCGACGCCGTCACCGA
CGGCAGCCTCCGGAGCCCGCTGCACCGCGAGTTACCAGGTCAACAGCGATTG

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GGGCAATGGCTTCACGGTAACGGTGGCCGTGACAAATTCCG-3' [SEQ ID NO.

3]

Translated amino acid sequence for W42R mutation, with modification
underlined.

5'-

AGGGYWHTSGREILDANNVPVRIAGINWFGFETCNVYVHGLRSRDYRSMLDQIK
SLGYNTIRLPYSDDILKPGTMPNSINFYQMNQDLQGLTSLQVMDKIVAYAGQIGL
RIILDRHRPDCSGQSALWYTSSVSEATWISDLQALAQRYKGNPTVVGFDLHNEPH
DPACWGC GDPSIDWRLAAERAGNAVLSVNP~~NLL~~IFVEGVQSYNGDSYWWGGN
LQGAGQYPVVLNVPNRLVYSAHDYATSVYPQTWFS DPTFPNNMPGIWNKNWG
YLFNQNIAPVWLGEFGTTLQSTTDQTLVQLRPTAQYGADSFQWTFWSW
NPDSGDTGGILKDDWQTVDTVKDGYLA~~PIK~~SSIFDPV-3' [SEQ ID NO. 3]

DNA sequence for Y82R Mutant with mutation site underlined.

5'-

GCGGGCGGCGGCTATTGGCACACGAGCGGCCGGGAGATCCTGGACGCGAAC
AACGTGCCGGTACGGATCGCCGGCATCAACTGGTTTGGGTTTCGAAACCTGCA
ATTACGTCGTGCACGGTCTCTGGTCACGCGACTACCGCAGCATGCTCGACCA
GATAAAGTCGCTCGGCTACAACACAATCCGGCTGCCGTACTCTGACGACATT
CTCAAGCCGGGCACCATGCCGAACAGCATCAATTTTCGGCAGATGAATCAGG
ACCTGCAGGGTCTGACGTCCTTGACAGGTCATGGACAAAATCGTCGCGTACGC
CGGTCAGATCGGCCTGCGCATCATTCTTGACCGCCACCGACCGGATTGCAGC
GGGCAGTCGGCGCTGTGGTACACGAGCAGCGTCTCGGAGGCTACGTGGATT
CCGACCTGCAAGCGCTGGCGCAGCGCTACAAGGGAAACCCGACGGTCGTCG
GCTTTGACTTGACAACGAGCCGCATGACCCGGCCTGCTGGGGCTGCGGCGA
TCCGAGCATCGACTGGCGATTGGCCGCCGAGCGGGCCGGAAACGCCGTGCTC
TCGGTGAATCCGAACCTGCTCATTTTCGTCGAAGGTGTGCAGAGCTACAACG
GAGACTCCTACTGGTGGGGCGGCAACCTGCAAGGAGCCGGCCAGTACCCGGT
CGTGCTGAACGTGCCGAACCGCCTGGTGTACTCGGCGCACGACTACGCGACG
AGCGTCTACCCGCAGACGTGGTTCAGCGATCCGACCTTCCCCAACACATGC
CCGGCATCTGGAACAAGAACTGGGGATACCTCTTCAATCAGAACATTGCACC

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GGTATGGCTGGGCGAATTCGGTACGACACTGCAATCCACGACCGACCAGACG
TGGCTGAAGACGCTCGTCCAGTACCTACGGCCGACCGCGCAATACGGTGCGG
ACAGCTTCCAGTGGACCTTCTGGTCCTGGAACCCCGATTCCGGGCGACACAGG
AGGAATTCTCAAGGATGACTGGCAGACGGTCGACACAGTAAAAGACGGCTAT
CTCGCGCCGATCAAGTCGTTCGATTTTCGATCCTGTCTAATGAATCGCCTAGCA
GTCAACCGTCCCCGTCGGTGTGCGCGTCTCCGTCGCCGAGCCCGTCGGCGAGT
CGGACGCCGACGCCTACTCCGACGCCGACAGCCAGCCCGACGCCAACGCTGA
CCCCTACTGCTACGCCACGCCACGGCAAGCCCGACGCCGTCACCGACGGC
AGCCTCCGGAGCCCGCTGCACCGCGAGTTACCAGGTCAACAGCGATTGGGGC
AATGGCTTCACGGTAACGGTGGCCGTGACAAATTCCG-3' [SEQ ID NO. 4]

Translated amino acid sequence for Y82R mutation, with modification
underlined.

5'-

AGGGYWHTSGREILDANNVPVRIAGINWFGFETCNVYVHGLWSRDYRSM LDQI
KSLGYNTIRLPYSDDILKPGTMPNSINFRQMNQDLQGLTSLQVMDKIVAYAGQIG
LRILDRHRPDCSGQSALWYTSSVSEATWISDLQALAQRYKGNPTVVGF DLHNEP
HDPACWGCGDPSIDWRLAAERAGNAVLSVNP NLLIFVEGVQSYNGDSYW
WGGNLQGAGQYPVVLNVPNRLVYSAHDYATSVYPQTWFS DPTFPNNMPGIWN
KNWGYLFNQNLAPVWLGEFGTTLQSTTDQTWL KTLVQYLRPTAQYGADSFQWT
FWSWNPDSGDTGGILKDDWQTVDTV KDGYLAPIKSSIFDPV-3' [SEQ ID NO. 4]